

# WEST Search History

DATE: Thursday, July 10, 2003

Set Name	Query	Hit Count	Set Name					
side by side	·		result set					
DB=USPT; PLUR=YES; OP=AND								
L1	nucleic or nucleotide or nuclear or dna or polynucleotide or cdna or mrna or rna	143841	L1					
L2	tag\$ or tagged or reporter\$2 or lable\$3 or label\$ or detect\$ or detection or detector\$ or marker\$2 or tracer\$2 or trace\$	1000254	L2					
L3	L2 near5 11	35360	L3					
L4	(protein\$ or peptide\$ or antigen\$ or analyte\$ or macromolecule or molecule or substance or target\$)	751144	L4					
L5	L3 same (ligase or polymerase or conjugate)	10278	L5					
L6	L3 near5 (ligase or polymerase or conjugate or amplification or amplify or amplifying)	6989	L6					
L7	L6 same 14	3814	L7					
L8	(detect\$ or measur\$ or determin\$ or diagnos\$ or test\$ or screen\$)	2080533	L8					
L9	11 same 12 same 14 same 18	30357	L9					
L10	11 near5 12 near5 14 near5 18	7720	L10					
L11	L10 same ligase	301	L11					
L12	(19 or 110)	30357	L12					
L13	L12 same (immunoassay or immunoper or antibody or monoclonal or mab or moab or atce)	12852	L13					
L14	L12 same (immunoassay or immunoper or antibody or monoclonal or mab or moab)	12558	L14					



	L15	L12 same (immunoassay or immunopcr)	2686	L15	
	L16	tag\$ or tagged or reporter\$2 or lable\$3 or label\$ or detector\$ or marker\$2 or trace\$	635016	L16	
	L17	L16 near5  1	28379	L17	
	L18	L17 same 18	20500	L18	
	L19	L16 near2  1	20324	L19	
	L20	L19 same 18 same 14	7006	L20	
	L21	L20 same (three or third or multiple) same (antibody or immunoassay or immunoper)	135	L21	
	DB=USI	PT,PGPB; PLUR=YES; OP=AND			
	L22	(4668621  4882269  5424413  5648213  5665539  5985548  6117631)![pn]	7	L22	
DB=USPT; PLUR=YES; OP=AND					
	L23	4957858.pn.	1	L23	
	L24	121 same ligat\$	6	L24	
	L25	121 same ligas\$	3	L25	
	L26	L25 or 124	9	L26	
	L27	epitope near2 proxim\$	98	L27	
	L28	L27 same (antibody or immunoassay or monoclonal or mab or moab or immunoper)	59	L28	
	L29	L28 same first same second	1	L29	
	L30	first same second same third same antibodies	2128	L30	
	L31	L30 same 11	443	L31	
	L32	L31 same 12	229	L32	
	L33	three near5 eptiope	0	L33	
	L34	three near5 epitope	797	L34	
	L35	L34 same 13	17	L35	



L36	134 same (antibody or immunoassay or monoclonal or mab or moab or immunopcr)	487	L36
L37	134 same 13	17	L37
L38	L34 same immobiliz\$	6	L38
L39	L34 same first same second same third	12	L39
L40	mutation\$ near10 epitope\$ near25 ligase	0	L40
L41	(mutation\$ near10 epitope\$) same ligase	1	L41
L42	(mutation\$ near10 epitope\$) same antibodies	152	L42
L43	142 same immunoassay	14	L43
L44	first.clm. and second.clm. and third.clm. and antibod\$.clm.	712	L44
L45	L44 and kit.clm.	187	L45
L46	(third or three).clm. same kit.clm.	979	L46
L47	L46 same lectin.clm.	0	L47
L48	L46 same 13	13	L48
L49	dna near binding near protein	3074	L49
L50	L49 same immunoassay	38	L50
L51	L49 same antibodies	417	L51
L52	L49 naser25 antibodies	0	L52
L53	L49 near25 antibodies	213	L53

END OF SEARCH HISTORY



U.S. Patent Dec. 11, 2001 Sheet 6 of 43 US 6,329,150 B1

## REPORTER ANTIBODIES

#### ASSAY

FIG.

# WEST

L21: Entry 45 of 135

File: USPT

Dec 11, 2001

DOCUMENT-IDENTIFIER: US 6329150 B1

TITLE: Unimolecular segment amplification and sequencing

#### Brief Summary Text (28):

Also disclosed are compositions and a method for of multiplex <u>detection of molecules</u> of interest involving rolling circle replication. The method is useful for simultaneously <u>detecting multiple</u> specific nucleic acids in a sample with high specificity and sensitivity. The method also has an inherently low level of background signal. A preferred form of the method consists of an association operation, an amplification operation, and a <u>detection</u> operation. The association operation involves association of one or more specially designed probe <u>molecules</u>, either wholly or partly nucleic acid, to <u>target molecules</u> of interest. This operation associates the probe <u>molecules</u> to a <u>target molecules</u> present in a sample. The amplification operation is rolling circle replication of circular nucleic acid <u>molecules</u>, termed amplification <u>target</u> circles, that are either a part of, or hybridized to, the probe <u>molecules</u>. A single round of amplification using rolling circle replication results in a large amplification of the amplification <u>target</u> circles, orders of magnitude greater than a single cycle of PCR replication and other amplification techniques in which each cycle is limited to a doubling of the number of copies of a <u>target</u> sequence. By coupling a <u>nucleic acid tag</u> to a specific binding <u>molecule</u>, such as an antibody, amplification of the <u>nucleic acid tag</u> can be used to <u>detect analytes</u> in a sample.

## WEST

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L21: Entry 70 of 135

File: USPT

Aug 29, 2000

DOCUMENT-IDENTIFIER: US 6110687 A

\*\* See image for Certificate of Correction \*\*

TITLE: Detection of antigens via oligonucleotide antibody conjugates

### **Brief Summary Text** (12):

The procedural complexity of immuno-PCR has been reduced by the direct chemical attachment of DNA to <u>analyte antibodies</u>, whereby immobilized capture <u>antibodies</u> and a reporter <u>antibody</u> that carries a covalently attached <u>DNA label</u> are used, and the assay response is obtained by PCR of the <u>DNA label and detection</u> of the <u>amplification products</u>. This technique has been modified to develop an immuno-PCR sandwich assay for <u>multiple analytes</u> (see R. D. Joerger, et al., Clinical Chemistry, 1995, 41 (9): 1371-1377; E. R. Hendrickson, et al., Nucl. Acids Res., 1995, 23 (3): 522-529; and T. Sano, et al., Science, 1992, 258: 120-122).